

```

chain nodes :
18 19 20 21 22 29 30 31 32
ring nodes :
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 23 24 25 26 27 28
chain bonds :
1-18 4-31 10-19 13-32 15-21 16-24 20-21 21-22 25-29 28-30
ring bonds :
1-2 1-6 2-3 3-4 4-5 4-7 5-6 5-10 7-8 7-11 8-9 8-14 9-10 11-12 12-13
13-14 13-15 14-17 15-16 16-17 23-24 23-28 24-25 25-26 26-27 27-28
exact/norm bonds :
1-2 1-6 1-18 2-3 3-4 4-5 4-7 5-6 5-10 7-8 7-11 8-9 8-14 9-10 10-19
11-12 12-13 13-14 13-15 14-17 15-16 16-17 21-22
exact bonds :
4-31 13-32 15-21 16-24 20-21 25-29 28-30
normalized bonds :
23-24 23-28 24-25 25-26 26-27 27-28

```

```

Match level :
1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS
10:CLASS 11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS
18:CLASS 19:CLASS 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS
26:CLASS 27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS

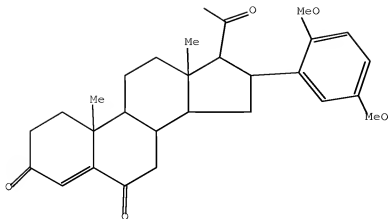
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L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l1 sam

SAMPLE SEARCH INITIATED 15:45:24 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 5 TO ITERATE

100.0% PROCESSED 5 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 5 TO 234
PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s l1 sss full

FULL SEARCH INITIATED 15:45:30 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 118 TO ITERATE

100.0% PROCESSED 118 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

L3 0 SEA SSS FUL L1

=> logoff hold

(FILE 'HOME' ENTERED AT 15:44:48 ON 12 MAY 2009)

FILE 'REGISTRY' ENTERED AT 15:45:02 ON 12 MAY 2009

L1 STRUCTURE UPLOADED

D L1

L2 0 SEA FILE=REGISTRY SSS SAM L1

L3 0 SEA FILE=REGISTRY SSS FUL L1

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
	185.88	186.10

FULL ESTIMATED COST

SESSION WILL BE HELD FOR 120 MINUTES

STN INTERNATIONAL SESSION SUSPENDED AT 15:45:44 ON 12 MAY 2009

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chain nodes :
18 19 20 21 28 29 30 31 38 39 40 41 42 43
ring nodes :
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 22 23 24 25 26 27
32 33 34 35 36 37
chain bonds :
1-43 4-30 10-18 13-31 15-20 19-20 20-21 24-28 27-29 32-40 33-38 36-42
37-41 38-39
ring bonds :
1-2 1-6 2-3 3-4 4-5 4-7 5-6 5-10 7-8 7-11 8-9 8-14 9-10 11-12 12-13
13-14 13-15 14-17 15-16 16-17 22-23 22-27 23-24 24-25 25-26 26-27 32-33
32-37 33-34 34-35 35-36 36-37
exact/norm bonds :
1-2 1-6 1-43 2-3 3-4 4-5 4-7 5-6 5-10 7-8 7-11 8-9 8-14 9-10 10-18
11-12 12-13 13-14 13-15 14-17 15-16 16-17 20-21 32-33 32-37 32-40 33-34
34-35 35-36 36-37 36-42 37-41 38-39
exact bonds :
4-30 13-31 15-20 19-20 24-28 27-29 33-38
normalized bonds :
22-23 22-27 23-24 24-25 25-26 26-27

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Match level :

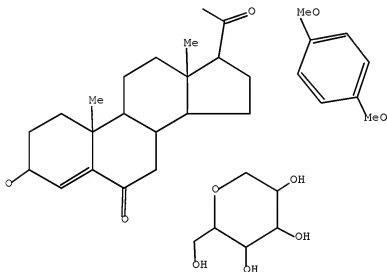
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18:CLASS 19:CLASS 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS
26:CLASS 27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS
34:CLASS 35:CLASS 36:CLASS 37:CLASS 38:CLASS 39:CLASS 40:CLASS 41:CLASS
42:CLASS 43:CLASS

L4 STRUCTURE UPLOADED

=> d l4

L4 HAS NO ANSWERS

L4 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l4 sss sam

SAMPLE SEARCH INITIATED 15:50:06 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 28 TO ITERATE

100.0% PROCESSED 28 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 243 TO 877

PROJECTED ANSWERS: 0 TO 0

L5 0 SEA SSS SAM L4

=> s l4 sss full

FULL SEARCH INITIATED 15:50:13 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 578 TO ITERATE

100.0% PROCESSED 578 ITERATIONS
SEARCH TIME: 00.00.01

0 ANSWERS

L6 0 SEA SSS FUL L4

=> file uspatfull

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	372.24	372.46

FILE 'USPATFULL' ENTERED AT 15:50:27 ON 12 MAY 2009
CA INDEXING COPYRIGHT (C) 2009 AMERICAN CHEMICAL SOCIETY (ACS)

FILE COVERS 1971 TO PATENT PUBLICATION DATE: 12 May 2009 (20090512/PD)
FILE LAST UPDATED: 12 May 2009 (20090512/ED)
HIGHEST GRANTED PATENT NUMBER: US7533422
HIGHEST APPLICATION PUBLICATION NUMBER: US20090119816
CA INDEXING IS CURRENT THROUGH 12 May 2009 (20090512/UPCA)
ISSUE CLASS FIELDS (/INCL) CURRENT THROUGH: 12 May 2009 (20090512/PD)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Feb 2009
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Feb 2009

USPATFULL now includes complete International Patent Classification (IPC)
reclassification data for the third quarter of 2008.

=> s us20060149045/pn
L7 1 US20060149045/PN

=> file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	1.53	373.99

FILE 'REGISTRY' ENTERED AT 15:51:09 ON 12 MAY 2009
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provided by InfoChem.

STRUCTURE FILE UPDATES: 10 MAY 2009 HIGHEST RN 1145355-82-3
DICTIONARY FILE UPDATES: 10 MAY 2009 HIGHEST RN 1145355-82-3

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH January 9, 2009.

Please note that search-term pricing does apply when
conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and
predicted properties as well as tags indicating availability of
experimental property data in the original document. For information
on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

=> tra 17 1- rn

L8 TRANSFER L7 1- RN : 386 TERMS
L9 386 L8

=> file caplus
COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
0.48	391.08

FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 15:51:37 ON 12 MAY 2009
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FILE COVERS 1907 - 12 May 2009 VOL 150 ISS 20
FILE LAST UPDATED: 10 May 2009 (20090510/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Feb 2009
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Feb 2009

Caplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

CAS Information Use Policies apply and are available at:

<http://www.cas.org/legal/infopolicy.html>

This file contains CAS Registry Numbers for easy and accurate

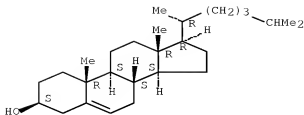
=> s l9 and cellobiosyl
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 281 CELLOBIOSYL
L10 5 L9 AND CELLOBIOSYL

=> d ibib ab hitstr 1-5

L10 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2007:1364352 CAPLUS Full-text
DOCUMENT NUMBER: 148:32596
TITLE: Nutraceutical compositions from microalgae and related
 methods of production and administration
INVENTOR(S): Dillon, Harrison F.; Somanchi, Aravind; Rao, Kamallesh;
 Jones, Peter J. H.
PATENT ASSIGNEE(S): Solazyme, Inc., USA
SOURCE: PCT Int. Appl., 199pp.
 CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 8
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007136428	A2	20071129	WO 2007-US1319	20070119
WO 2007136428	A3	20081127		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA			
US 20070167396	A1	20070719	US 2006-336428	20060119
US 20070167397	A1	20070719	US 2006-336430	20060119
US 20070166449	A1	20070719	US 2006-336431	20060119
US 20070166797	A1	20070719	US 2006-336656	20060119
US 20070166266	A1	20070719	US 2006-337103	20060119
US 20070167398	A1	20070719	US 2006-337171	20060119
US 20070191303	A1	20070816	US 2006-336426	20060119
EP 1993565	A2	20081126	EP 2007-808975	20070119
R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS			
PRIORITY APPLN. INFO.:			US 2006-336426	A 20060119
			US 2006-336428	A 20060119
			US 2006-336430	A 20060119
			US 2006-336431	A 20060119
			US 2006-336656	A 20060119
			US 2006-337103	A 20060119
			US 2006-337171	A 20060119
			US 2006-816967P	P 20060628
			US 2006-832091P	P 20060720
			US 2006-838452P	P 20060817
			US 2006-872072P	P 20061130
			WO 2007-US1319	W 20070119
AB	Polysaccharides with nutraceutical application may be obtained by culturing red microalgae and the nutraceutical compns. thus produced may comprise a carrier and homogenized microalgal cells. Addnl. components may include phytosterols, limonoids, flavonoids, and tocotrienols. The polysaccharides may be used in applications such as reducing cholesterol in mammals, inactivating viruses, stabilizing foods, etc. Thus, total serum cholesterol in an animal model (hamsters) over 30 days was decreased 35-62% by dietary inclusion of Porphyridium biomass homogenate and polysaccharide, the highest decreases being observed when phytosterols were also present. Transgenic algae may be used that are capable of utilizing fixed carbon sources for energy. Also provided are novel nucleic acid sequences from red microalgae.			
IT	57-88-5, Cholest-5-en-3-ol (3 β)-, biological studies RL: BSU (Biological study, unclassified); BIOL (Biological study) (blood; nutraceutical compns. from red microalgae and related methods of production and administration)			
RN	57-88-5 CAPLUS			
CN	Cholest-5-en-3-ol (3 β)- (CA INDEX NAME)			

Absolute stereochemistry.



L10 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2000:490669 CAPLUS [Full-text](#)

DOCUMENT NUMBER: 133:250148

TITLE: Intestinal absorption of cholesterol is mediated by a saturable, inhibitable transporter

AUTHOR(S): Hernandez, M.; Montenegro, J.; Steiner, M.; Kim, D.; Sparrow, C.; Detmers, P. A.; Wright, S. D.; Chao, Y.-S.

CORPORATE SOURCE: Merck Research Laboratories, Rahway, NJ, 07065, USA

SOURCE: Biochimica et Biophysica Acta, Molecular and Cell Biology of Lipids (2000), 1486(2-3), 232-242
CODEN: BBMLFG; ISSN: 1388-1981

PUBLISHER: Elsevier B.V.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Although the mechanism by which dietary cholesterol is absorbed from the intestine is poorly understood, it is generally accepted that cholesterol is absorbed from bile acid micelles in the jejunum. Once inside the enterocytes, cholesterol is esterified by the action of acyl-CoA:cholesterol acyltransferase (ACAT), assembled into chylomicrons, and secreted into the lymph. In this work, mechanistic aspects of cholesterol absorption were probed using compds. that block cholesterol absorption in hamsters. Sterol glycoside cholesterol absorption inhibitors, exemplified by L-166,143, (3 β ,5 α ,25R)-3-[(4'',6''-bis[2-fluoro-phenylcarbamoyl]-B-D- cellobiosyl)oxy]-spirostan-11-one, potentially blocked absorption of radioactive cholesterol, and the potencies of several analogs correlated with their ability to lower plasma cholesterol. Each mol. of L-166,143 blocked the uptake of 500 mols. of cholesterol, rendering it unlikely that the inhibitor interacts directly with the cholesterol or bile acid. Radiolabeled L-166,143 bound to the mucosa and binding was blocked by active, but not inactive, cholesterol absorption inhibitors. Subtle changes in the structure of sterol glycosides yielded large changes in their ability to block both cholesterol absorption and binding of radiolabeled L-166,143. dog. Large species-to-species variation in potency was also observed. These lines of evidence support the interpretation that dietary cholesterol is absorbed via a specific transporter found in the intestinal mucosa.

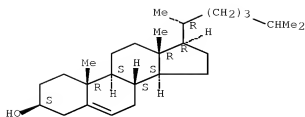
IT 57-88-5, Cholesterol, biological studies

RL: BAC (Biological activity or effector, except adverse); BOC (Biological occurrence); BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); OCCU (Occurrence); PROC (Process) (intestinal absorption of cholesterol mediated by saturable inhibitable transporter in hamsters, dogs, rats and mice)

RN 57-88-5 CAPLUS

CN Cholest-5-en-3-ol (3 β)- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2009 ACS on STN
 ACCESSION NUMBER: 1994:509562 CAPLUS Full-text
 DOCUMENT NUMBER: 121:109562
 ORIGINAL REFERENCE NO.: 121:19815a,19818a
 TITLE: Steroidal glycosides for treating hypercholesterolemia
 INVENTOR(S): Deninno, Michael Paul; McCarthy, Peter Andrew
 PATENT ASSIGNEE(S): Pfizer Inc., USA
 SOURCE: PCT Int. Appl., 85 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

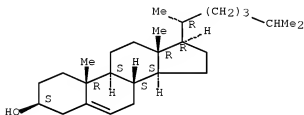
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9400480	A1	19940106	WO 1993-US4092	19930506
W: AU, BG, BR, CA, CZ, DE, JP, KR, NO, NZ, RO, RU, SK, UA, US				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
AU 9342265	A	19940124	AU 1993-42265	19930506
EP 647234	A1	19950412	EP 1993-910951	19930506
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE				
JP 07504921	T	19950601	JP 1993-502331	19930506
EP 796862	A2	19970924	EP 1997-200454	19930506
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE				
EP 796863	A2	19970924	EP 1997-200455	19930506
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE				
EP 796864	A2	19970924	EP 1997-200456	19930506
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE				
BR 9306619	A	19981208	BR 1993-6619	19930506
CN 1085561	A	19940420	CN 1993-107620	19930625
ES 2074006	A1	19950816	ES 1993-1507	19930705
ES 2074006	B1	19960316		
US 5629295	A	19970513	US 1994-351470	19941220
NO 9405001	A	19950214	NO 1994-5001	19941223
JP 09309897	A	19971202	JP 1997-30588	19941226
PRIORITY APPLN. INFO.:				
US 1992-904914 A2 19920626				
EP 1993-910951 A3 19930506				
WO 1993-US4092 A 19930506				
JP 1994-502331 A3 19941226				

OTHER SOURCE(S): MARPAT 121:109562
 AB Certain steroidal glycosides of formula I [e.g., Q1 = CO, CH(OH); Q2 = CO, CH2, CH(OH); Q3 = CH(OR1), CH(OXOR1); Q4, Q5 = CH2; R1 = various glycosyl residues; X = alkylene; plus several addnl. groups of definitions], useful as

hypocholesterolemic and antiatherosclerotic agents (no data), are claimed and prepared For example, ZnF2-promoted coupling of (3 β ,5 α ,25R)-3-hydroxyspirostan-11-one with heptaacetyl- β -D-cellobiosyl bromide (93% yield) and deacetylation with NaOMe in MeOH-THF (57% yield) gave the invention compound (3 β ,5 α ,25R)-3-[(β -D-cellobiosyl)oxy]spirostan-11-one. Prepn. of approx. 50 I and numerous precursors are described.

IT 57-88-5, Cholesterol, biological studies
 RL: BIOL (Biological study)
 (absorption of, inhibitors of, steroidal glycosides as)
 RN 57-88-5 CAPLUS
 CN Cholest-5-en-3-ol (3 β)- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1990:50901 CAPLUS Full-text

DOCUMENT NUMBER: 112:50901

ORIGINAL REFERENCE NO.: 112:8649a,8652a

TITLE: ESR study on synthetic glyceroglycolipid liposomal membranes

AUTHOR(S): Naito, Mikihiro; Utsumi, Hideo; Umeda, Masato; Kudo, Ichiro; Takeshita, Keizo; Hamada, Akira; Nojima, Shoshichi; Inoue, Keizo

CORPORATE SOURCE: Fac. Pharm. Sci., Univ. Tokyo, Tokyo, 113, Japan
 SOURCE: Biochimica et Biophysica Acta, Biomembranes (1989), 985(2), 147-52

CODEN: BBBMBS; ISSN: 0005-2736

DOCUMENT TYPE: Journal

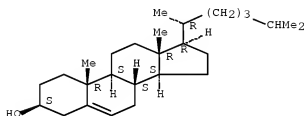
LANGUAGE: English

AB It was previously reported that glyceroglycolipid liposomes without cholesterol activated mouse peritoneal macrophages in vivo and in vitro, whereas glyceroglycolipid liposomes containing equimolar cholesterol did not. In order to characterize the properties of the glyceroglycolipid membranes, ESR spectroscopic studies were carried out with an acyl spin-labeled galactosyl ceramide (SL-GC) or a headgroup spin-labeled phospholipid (SL-6-DPPA) in 1,2-dipalmityl[β -cellobiosyl (1'→3)]glycerol (Cel-DAG) liposomal membranes. The ESR spectrum of the SL-GC in the Cel-DAG liposomes at 37° was a single broad line, indicating that the SL-GC mols. were excluded almost completely from Cel-DAG domains and formed clusters in the membranes. The spectrum of SL-6-DPPA in the Cel-DAG liposomes at 37° showed broad resonance lines with the central peak being the highest, while that at 60° gave narrow lines with the low-field peak being the highest. This observation and rotational correlation time anal. showed that the mol. motions of the spin-label moiety of the SL-6-DPPA were extremely restricted at 37°C but not above

Tc. These results suggest that below Tc the Cel-DAG mols. are packed tightly and restricted in motion in the membrane. Incorporation of cholesterol into the Cel-DAG liposomal membranes gave (1) the spectra of the SL-GC triplet, and (2) the spectra of the SL-6-DPPA narrow resonance with the low-field peak being the highest. Apparently, cholesterol disturbs the rigid-packed structure of the Cel-DAG membrane and increases the mol. motions of the Cel-DAG. The DSC anal. of Cel-DAG with and without cholesterol agreed well with the results of the ESR technique. Thus it is assumed that peritoneal macrophages recognize the rigid-packed carbohydrate residues which are restricted in motion on the Cel-DAG membranes.

IT 57-88-5, Cholesterol, biological studies
 RL: BIOL (Biological study)
 (glyceroglycolipid membrane fluidity response to)
 RN 57-88-5 CAPLUS
 CN Cholest-5-en-3-ol (3 β)- (CA INDEX NAME)

Absolute stereochemistry.



L10 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1987:475887 CAPLUS [Full-text](#)

DOCUMENT NUMBER: 107:75887

ORIGINAL REFERENCE NO.: 107:12489a,12492a

TITLE: Activation of mouse peritoneal macrophages by
 synthetic glyceroglycolipid liposomes
 AUTHOR(S): Naito, Mikihiko; Kudo, Ichiro; Mukai-Sato, Yukiko;
 Tsushima, Susumu; Nomura, Hiroaki; Nojima, Shoshichi;
 Inoue, Keizo

CORPORATE SOURCE: Fac. Pharm. Sci., Univ. Tokyo, Tokyo, 113, Japan
 SOURCE: Cancer Immunology Immunotherapy (1987), 24(2), 158-64
 CODEN: CIIMDN; ISSN: 0340-7004

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Liposomes composed of chem. synthesized glyceroglycolipids, such as 1,2-dipalmityl-[β -cellobiosyl-(1'→3)]-glycerol (Cel-DAG), 1,2-dipalmityl-[β -lactosyl-(1'→3)]-glycerol, or 1,2-dipalmityl-[β -maltosyl-(1'→3)]-glycerol, enhanced protective immunity against transplantable tumor cells (sarcoma 180) in ICR mice. Peritoneal exudate cells prepared from mice treated in vivo with Cel-DAG showed cytostatic activity in vitro against the mouse leukemia cell line, EL-4. Adherent cells separated from this preparation showed similar activity. Peritoneal cells from polypeptide-injected mice acquired appreciable cytostatic activity when incubated in vitro in the presence of glyceroglycolipid liposomes. The adherent cell fraction alone showed rather weak cytostatic activity when pretreated with the glyceroglycolipids, and full activity was restored by supplementing with the nonadherent cell fraction. The ability of glycolipids to induce tumoricidal effects was affected by cholesterol content: with increasing cholesterol content, the activities decreased. Cholesterol-free glycolipid liposomes were taken more efficiently

by macrophages than cholesterol-containing liposomes. Cholesterol modifies the surface property of glyceroglycolipid liposomes. Activation of macrophages is responsible for enhancement of protective immunity against tumor cells by injection of these glycolipids in vivo.

IT 57-88-5, Cholesterol, biological studies

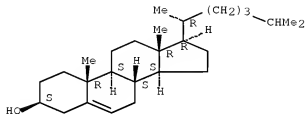
RL: BIOL (Biological study)

(glyceroglycolipid liposomes activation of macrophages modulation by)

RN 57-88-5 CAPLUS

CN Cholest-5-en-3-ol (3 β)- (CA INDEX NAME)

Absolute stereochemistry.



=> d his

(FILE 'HOME' ENTERED AT 15:44:48 ON 12 MAY 2009)

FILE 'REGISTRY' ENTERED AT 15:45:02 ON 12 MAY 2009

L1 STRUCTURE UPLOADED
L2 0 S L1 SAM
L3 0 S L1 SSS FULL
L4 STRUCTURE UPLOADED
L5 0 S L4 SSS SAM
L6 0 S L4 SSS FULL

FILE 'USPATFULL' ENTERED AT 15:50:27 ON 12 MAY 2009
1 S US20060149045/PN

FILE 'REGISTRY' ENTERED AT 15:51:09 ON 12 MAY 2009

FILE 'USPATFULL' ENTERED AT 15:51:17 ON 12 MAY 2009
L8 TRA L7 1- RN : 386 TERMS

FILE 'REGISTRY' ENTERED AT 15:51:18 ON 12 MAY 2009
L9 386 SEA L8

FILE 'CAPLUS' ENTERED AT 15:51:37 ON 12 MAY 2009
L10 5 S L9 AND CELLOBIOSYL

=> logoff hold

(FILE 'HOME' ENTERED AT 15:44:48 ON 12 MAY 2009)

FILE 'REGISTRY' ENTERED AT 15:45:02 ON 12 MAY 2009

L1 STRUCTURE UPLOADED
D L1
L2 0 SEA SSS SAM L1

L3 0 SEA SSS FUL L1
L4 STRUCTURE UPLOADED
D L4
L5 0 SEA SSS SAM L4
L6 0 SEA SSS FUL L4

FILE 'USPATFULL' ENTERED AT 15:50:27 ON 12 MAY 2009
L7 1 SEA SPE=ON ABB=ON PLU=ON US20060149045/PN

FILE 'REGISTRY' ENTERED AT 15:51:09 ON 12 MAY 2009

FILE 'USPATFULL' ENTERED AT 15:51:17 ON 12 MAY 2009
L8 TRA PLU=ON L7 1- RN : 386 TERMS

FILE 'REGISTRY' ENTERED AT 15:51:18 ON 12 MAY 2009
L9 386 SEA SPE=ON ABB=ON PLU=ON L8

FILE 'CAPLUS' ENTERED AT 15:51:37 ON 12 MAY 2009
L10 5 SEA SPE=ON ABB=ON PLU=ON L9 AND CELLOBIOSYL
D IBIB AB HITSTR 1-5

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	31.94	423.02
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-4.10	-4.10

SESSION WILL BE HELD FOR 120 MINUTES
STN INTERNATIONAL SESSION SUSPENDED AT 15:53:11 ON 12 MAY 2009

6.3° (c 0.526, CHCl₃). I maltoside, fine needles, m. 288°, [α]_D¹⁷ 22.3° (c 0.735, CHCl₃). In the same way, 500 mg. β'-(Δ⁵-3β-hydroxynorcholen-23-yl)-Δα',β'-butenolide (IV) and acetobromoglucose give 385 mg. IV tetraacetylglucoside, fine needles, m. 208-8.5°, in addition to 300 mg. recovered IV; IV glucoside, prepared by saponification with 0.1 N Ba(OMe)₂ in MeOH at -10° for 2 days, crystals from EtOH, m. 270-5°. It gives a pos. L. test. Δ²⁰,22-3β,21-Dihydroxycholeonic acid lactone tetraacetylglucoside, crystal, from iso-AmOMe, m. 125-7°. These compds. have only a slight solubility in H₂O. All m.ps. corrected and in evacuated tube.

=> d his

(FILE 'HOME' ENTERED AT 00:48:14 ON 13 MAY 2009)

FILE 'REGISTRY' ENTERED AT 00:48:33 ON 13 MAY 2009

```
L1      STRUCTURE UPLOADED
L2      2 S L1 SAM
L3      65 S L1 SSS FULL
```

FILE 'CAPLUS' ENTERED AT 00:50:47 ON 13 MAY 2009

```
L4      6 S L3
L5      3839 S STEROID AND (GLUCO OR GLYCO OR CELLOBIOSYL OR CELLBIOSE OR
L6      3722 S L5 AND PY<2002
L7      190 S STEROID AND (CELLBIOSYL OR CELLOBIOSIDE OR GLYCOSYL OR GLYCOS
L8      0 S L7 AND 3-O-CELLOBIOSYL
L9      0 S L8 AND 3-O-CELLOBIOSIDE
        E CANCER+ALL/CT
L10     1 S L7 AND C16
L11     10 S STEROID AND (CELLBIOSYL OR CELLOBIOSIDE) AND PY<2002
```

=> logoff hold

(FILE 'HOME' ENTERED AT 00:48:14 ON 13 MAY 2009)

FILE 'REGISTRY' ENTERED AT 00:48:33 ON 13 MAY 2009

```
L1      STRUCTURE UPLOADED
        D L1
L2      2 SEA SSS SAM L1
L3      65 SEA SSS FUL L1
```

FILE 'CAPLUS' ENTERED AT 00:50:47 ON 13 MAY 2009

```
L4      6 SEA SPE=ON ABB=ON PLU=ON L3
        D IBIB AB HITSTR 1-6
L5      3839 SEA SPE=ON ABB=ON PLU=ON STEROID AND (GLUCO OR GLYCO OR
        CELLOBIOSYL OR CELLBIOSE OR GLYCOSYL OR GLYCOSYLATED)
L6      3722 SEA SPE=ON ABB=ON PLU=ON L5 AND PY<2002
L7      190 SEA SPE=ON ABB=ON PLU=ON STEROID AND (CELLBIOSYL OR
        CELLOBIOSIDE OR GLYCOSYL OR GLYCOSYLATED) AND PY<2002
L8      0 SEA SPE=ON ABB=ON PLU=ON L7 AND 3-O-CELLOBIOSYL
L9      0 SEA SPE=ON ABB=ON PLU=ON L8 AND 3-O-CELLOBIOSIDE
        SET LINE 250
        SET DETAIL OFF
        E CANCER+ALL/CT
        SET LINE LOGIN
        SET DETAIL LOGIN
L10     1 SEA SPE=ON ABB=ON PLU=ON L7 AND C16
        D IBIB
L11     10 SEA SPE=ON ABB=ON PLU=ON STEROID AND (CELLBIOSYL OR
        CELLOBIOSIDE) AND PY<2002
        D IBIB AB 1-10
```

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	131.09	318.63